

Title 21—Food and Drugs

(This book contains parts 300 to 499)

Part

CHAPTER I—Food and Drug Administration, Department of Health and Human Services (Continued)	300
--	-----

CROSS REFERENCES: Food Safety and Inspection Service, Department of Agriculture: See 9
CFR chapter III.

Federal Trade Commission: See Commercial Practices, 16 CFR chapter I.

United States Customs Service, Department of the Treasury: See Customs Duties, 19 CFR
chapter I.

Internal Revenue Service, Department of the Treasury: See Internal Revenue, 26 CFR chap-
ter I.

Bureau of Alcohol, Tobacco, and Firearms, Department of the Treasury: See Alcohol, To-
bacco Products and Firearms, 27 CFR chapter I.

CHAPTER I—FOOD AND DRUG ADMINISTRATION, DEPARTMENT OF HEALTH AND HUMAN SERVICES—Continued

(Parts 300 to 499)

EDITORIAL NOTE: The Food and Drug Administration published a document at 49 FR 41019, Oct. 19, 1984, establishing July 1, 1987, as a uniform effective date for compliance for regulations affecting the labeling of food products. At 51 FR 34085, Sept. 25, 1986, FDA established January 1, 1989, as a new uniform effective date for compliance. The new uniform effective date will apply only to final FDA food labeling regulations published after July 1, 1986, and before January 1, 1988. At 55 FR 276, Jan. 4, 1990, FDA established January 1, 1993 as a new uniform effective date for compliance. The new uniform effective date will apply only to final FDA food labeling regulations published after January 1, 1990 and before January 1, 1992.

SUBCHAPTER D—DRUGS FOR HUMAN USE

<i>Part</i>		<i>Page</i>
300	General	5
310	New drugs	5
312	Investigational new drug application	57
314	Applications for FDA approval to market a new drug	95
315	Diagnostic radiopharmaceuticals	172
316	Orphan drugs	174
320	Bioavailability and bioequivalence requirements ...	186
328	Over-the-counter drug products intended for oral ingestion that contain alcohol	200
329	Habit-forming drugs	202
330	Over-the-counter (OTC) human drugs which are generally recognized as safe and effective and not misbranded	206
331	Antacid products for over-the-counter (OTC) human use	219
332	Antiflatulent products for over-the-counter human use	222
333	Topical antimicrobial drug products for over-the- counter human use	224
336	Antiemetic drug products for over-the-counter human use	231

21 CFR Ch. I (4–1–01 Edition)

<i>Part</i>		<i>Page</i>
338	Nighttime sleep-aid drug products for over-the-counter human use	233
340	Stimulant drug products for over-the-counter human use	234
341	Cold, cough, allergy, bronchodilator, and anti-asthmatic drug products for over-the-counter human use	235
343	Internal analgesic, antipyretic, and antirheumatic drug products for over-the-counter human use	252
344	Topical OTIC drug products for over-the-counter human use	260
346	Anorectal drug products for over-the-counter human use	262
347	Skin protectant drug products for over-the-counter human use	267
348	External analgesic drug products for over-the-counter human use	269
349	Ophthalmic drug products for over-the-counter human use	270
352	Sunscreen drug products for over-the-counter human use	275
355	Anticaries drug products for over-the-counter human use	285
357	Miscellaneous internal drug products for over-the-counter human use	289
358	Miscellaneous external drug products for over-the-counter human use	293
361	Prescription drugs for human use generally recognized as safe and effective and not misbranded: Drugs used in research	300
369	Interpretative statements re warnings on drugs and devices for over-the-counter sale	305
370–499	[Reserved]	

EDITORIAL NOTE: For nomenclature changes to chapter I see 59 FR 14366, Mar. 28, 1994.